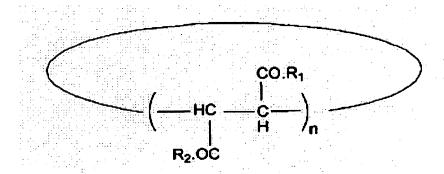
AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions and listings of claims in the application:

Claims 1-4 (Canceled)

5. (Currently amended) An carbocyclic fumaric acid oligomer according to claim 1 or 2 of the following formula (I)



wherein the radicals R_1 and R_2 are the same or different and <u>each occurrence of radicals R_1 and R_2 is independently [[are]] selected chosen from the group consisting of amine radicals (-NR₃R₄), amino acid radicals (-NH-CH(COOH)-R₆), peptide radicals having <u>from 2</u> to 100 amino acids, alcohol radicals (-OR₅) and a hydroxyl radical,</u>

n is an integer from between 2 and to 10,

the radicals R_3 and R_4 are the same or different and are <u>independently chosen</u> selected from the group consisting of hydrogen, C_{1-24} alkyl radicals, the <u>a</u> phenyl radical and C_{6-10} aralkyl radicals,

the radical R_5 is <u>chosen</u> selected from the group consisting of hydrogen, C_{1-24} alkyl radicals, the <u>a</u> phenyl radical and C_{6-10} aralkyl radicals,

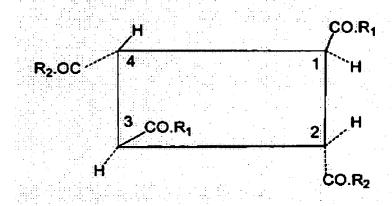
and the radical R_6 represents the side chain of a natural or synthetic amino acid, provided that when R_1 and R_2 are alcohol radicals (-OR₅) and R_5 is a methyl radical, then n is not 2, further provided that when R_1 and R_2 are hydroxyl radicals, then n is not 3, and even further provided that when R_1 is an amine radical (-NR₃R₄), R_3 is a hydrogen, R_4 is a C_{6-10} aralkyl radical, and R_2 is hydrogen, then n is not 2.

6. (Canceled)

- 7. (Currently amended) An oligomer according to claim 5 er-6, wherein the radicals R₃, R₄, and R₅ are the same or different and are <u>independently chosen</u> selected from the group consisting of hydrogen, methyl, ethyl, n-propyl, isopropyl, n-butyl, secbutyl, t-butyl, pentyl, cyclopentyl, 2-ethyl hexyl, hexyl, cyclohexyl, heptyl, cycloheptyl, octyl, vinyl, allyl, 2-hydroxyethyl, 2-hydroxypropyl, [[or]] 3-hydroxypropyl, 2,3-dihydroxypropyl, 2-methoxyethyl, methoxymethyl, and 2-methoxypropyl, and [[or]] 3-methoxypropyl.
- 8. (Currently amended) An oligomer according to claim 5 or 6 wherein R₁ represents an amine radical and <u>each occurrence of R₂ represents an is independently chosen from alkoxy alcohol radicals (-OR₅) and a hydroxyl radical [[or -OH]].</u>
- 9. (Currently amended) An oligomer according to claim 5 or 6 wherein each occurrence of R₁ and R₂ each independently is independently chosen from alcohol radicals (-OR₅) and represent an alkoxy radical or a hydroxyl radical.

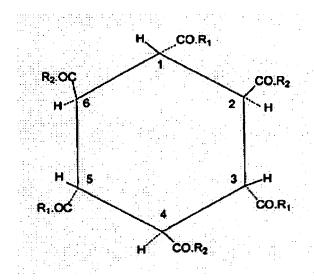
- 10. (Currently amended) An oligomer according to claim 9 wherein <u>each</u>

 <u>occurrence of</u> R₁ and R₂ [[are]] <u>is</u> independently <u>chosen</u> selected from the group
 consisting of methoxy, ethoxy, propoxy, isopropoxy, n-butoxy, sec-butoxy, tert-butoxy, phenoxy, and pentoxy.
- 11. (Original) An oligomer according to claim 9 wherein R₁ and R₂ are both methoxy.
- 12. (Currently amended) An earbocyclic or exacarbocyclic oligomer according to any of the previous claims claim 5, wherein n is 2 or 3, each occurrence of radicals R₁ and R₂ is independently chosen from alcohol radicals (-OR₅), and R₅ is a C₁₋₂₄ alkyl radical containing 2 to 3 dialkyl fumarate units derived from fumaric acids and/or esters and amides thereof.
- 13. (Currently amended) $A\underline{n}$ carbocyclic oligomer according to any of the previous claims claim 12 wherein all of the carbonyl groups carrying the radicals R_1 and R_2 are arranged as substituents in the trans position to each adjacent substituent.
- 14. (Currently amended) A<u>n</u>-carbocyclic oligomer according to claim 5 represented by the formula (II)



wherein R₁ and R₂ are as defined in claim 5.

15. (Currently amended) An earbocyclic oligomer according to claim 5 represented by the formula (III)



wherein R₁ and R₂ are as defined in claim 5.

Claims 16-19 (Canceled)

- 20. (Currently amended) A pharmaceutical preparation containing comprising an fumaric acid oligomer according to claim 5 any of the claims 1 to 17 and at least one excipient.
- 21. (Original) A pharmaceutical preparation according to claim 20, said pharmaceutical preparation being available in a form suitable for oral, rectal, transdermal, dermal, ophthalmological, nasal, pulmonary or parenteral application.
- 22. (Currently amended) A pharmaceutical preparation according to claim 20, said pharmaceutical preparation being present in the form of tablets, coated tablets, capsules, granulate, solutions for drinking, liposomes, nano-particles, nano-capsules, micro-capsules, micro-tablets, pellets, [[or]] powders, as well as granulate filled in capsules, micro-tablets filled in capsules, pellets filled in capsules, nano-particles filled in capsules or powder filled in capsules.
- 23. (Currently amended) A pharmaceutical preparation according to claim 22, said pharmaceutical preparation being present in the form of nano-particles, micropellets or micro-tablets which, optionally, may be filled in sachets or capsules.
- 24. (Currently amended) A pharmaceutical preparation according to claim 22 wherein the solid oral dosage forms are provided with further comprise an enteric coating.

- 25. (Currently amended) A pharmaceutical preparation according to any of the claims 20 to 24 which contains an amount of fumaric acid an oligomer corresponding to 10 to 500 mg of fumaric acid.
- 26. (Currently amended) A method The use according to any of the claims 19 to 20 for preparing a pharmaceutical preparation comprising admixing an oligomer according to claim 5 with at least one excipient.
- (1) for the therapy of an autoimmune disease selected from the group consisting of polyarthritis, multiple sclerosis, graft-versus-host reactions, juvenile-onset diabetes, Hashimoto's thyroiditis, Grave's disease (Basedow disease), systemic Lupus-erythematodes (SLE), Sjogren's syndrome, pernicious anaemia and chronic active (= lupoid) hepatitis;
 - (2) for use in transplantation medicine; and
- (3) for the therapy of mitochondrial diseases selected from the group consisting of Parkinson syndrome, Alzheimer's disease, Chorea Huntington disease, retinopathia-pigrnentosa or forms of mitochondrial encephalomyopathy; as well as (4) for the therapy of NF-kappaB mediated diseases selected from the group consisting of progressive-systemic sclerodermia, osteochondritis syphilitica (Wegener's disease), cutis marmorata (livedo reticularis), Behcet disease, panarteriitis, colitis ulcerosa, vasculitis, osteoarthritis, gout, arteriosclerosis, Reiter's disease, pulmonary granulomatosis, types of encephalitis, endotoxic shock (septic-toxic-shock), sepsis, pneumonia, encephalomyelitis, anorexia nervosa, hepatitis (acute hepatitis, chronic hepatitis, toxic-hepatitis, alcohol-induced hepatitis, viral hepatitis, jaundice, liver insufficiency and

cytomogaloviral hopatitis), Rennert T-lymphomatosis, mesangial nephritis, postangioplastic restenosis, reperfusion syndrome, cytomogaloviral retinopathy, adenoviral
diseases such as adenoviral colds, adenoviral pharyngoconjunctival fever andadenoviral ophthalmia, AIDS, Guillain-Barré syndrome, post-horpetic or post-zosterneuralgia, inflammatory demyelinising polyneuropathy, menoneuropathia multiplex,
mucoviscidosis, Bechterew's disease, Barett oesophagus, EBV (Epstein-Barr virus)
infection, cardiac remodeling, interstitial cystitis, diabetes mellitus type 11,
radiosensitisation of malignant tumours, multi-resistance of malignant cells tochemotherapeutic agents (multipharmaceutical preparation resistance inchemotherapy), granuloma annulare and cancers such as mamma carcinoma, coloncarcinoma, melanoma, primary liver cell carcinoma, adenocarcinoma, kaposi's sarcoma,
prostate carcinoma, leukaemia such as acute myeloid leukaemia, multiple myeloma(plasmocytoma), Burkitt lymphoma and Castleman tumour.

- 27. (New) A method of treating an autoimmune disease comprising administering a pharmaceutical preparation according to any one of claims 20 to 24.
- 28. (New) A method of treating host-versus-graft reactions comprising administering a pharmaceutical preparation according to any one of claims 20 to 24.
- 29. (New) A method for treating mitochondrial diseases comprising administering a pharmaceutical preparation according to any one of claims 20 to 24.

- 30. (New) A pharmaceutical preparation according to claim 23, wherein said nano-particles, micro-pellets or micro-tablets are filled in sachets or capsules.
- 31. (New) A method for preparing a pharmaceutical preparation according to claim 26 further comprising subjecting the admixture to tabletting, direct compression, melt methods, or spray drying to form tablets, granulates, nano-particles, nano-capsules, micro-capsules, micro-tablets, pellets, or powders.
- 32. (New) A method for preparing a pharmaceutical preparation according to claim 31, wherein said tablets, granulates, nano-particles, nano-capsules, micro-capsules, micro-tablets, pellets, or powders are enterically coated.
- 33. (New) A method for preparing a pharmaceutical preparation according to claim 31, wherein said nano-particles, nano-capsules, micro-capsules, micro-tablets, pellets, or powders are put into capsules.
- 34. (New) An oligomer according to claim 5, wherein n is 2 or 3, R_1 is hydrogen, R_2 is an alcohol radical (-OR₅), and R_5 is a C_{1-24} alkyl radical.
- 35. (New) An oligomer according to claim 5, wherein n is 3, R_1 is hydrogen, R_2 is an amine radical (-NR₃R₄).

- 36. (New) An oligomer according to claim 5, wherein n is 2 or 3, R_1 and R_2 are independently chosen from amine radicals (-NR₃R₄).
- 37. (New) An oligomer according to claim 5, wherein n is 2 or 3, R_1 is an alcohol radical (-OR₅), R_5 is a C_{1-24} alkyl radical, and R_2 is an amine radical (-NR₃R₄).